This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:** 

Claim 1 (currently amended) An antagonist of ghrelin, wherein the antagonist is a nucleic

acid which specifically binds, and whereby preferably the nucleic acid is binding to ghrelin.

Claim 2 (currently amended) An antagonist of the GHSR 1a receptor system, wherein the
antagonist is a which is the nucleic acid, and whereby preferably the nucleic acid is binding to

the ligand of the receptor and whereby preferably the ligand is which specifically binds to

ghrelin of claim 1.

Claim 3 (original) The antagonist according to claim 1 or 2, wherein the nucleic acid

comprises at least one L-nucleotide.

Claim 4 (currently amended) The antagonist according to any of claim[[s]] 1 or 2 to 3,

wherein the antagonist is an L-nucleic acid.

Claim 5 (canceled)

Claim 6 (currently amended) A nucleic acid binding to ghrelin, preferably a L nucleic

acid binding which specifically binds to L-ghrelin.

Claim 7 (canceled)

Claim 8 (currently amended) A nucleic acid having a sequence which is selected from the

group comprising the sequences according to SEQ. ID. No. 7 to SEQ. ID. No. 125 which is one

of SEQ ID NO:7 to SEQ ID NO:125.

Claim 9 (original) The nucleic acid according to claim 8, wherein the nucleic acid

comprises at least one L-nucleotide.

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Claim 10 (currently amended) The nucleic acid according to any of claims 8 to claim 8 or 9, wherein the nucleic acid is an L-nucleic acid.

Claim 11 (currently amended) The nucleic acid according to any of claims 8 to 10, of

claim 8 or 9 wherein the nucleic acid is selected from the group comprising consisting of DNA,

RNA and combinations thereof.

Claim 12 (currently amended) The nucleic acid according to any of claims 8 of claim 8

 $\underline{\text{or 9}},$  to 11 wherein the Kd of the nucleic acid is less than 1  $\mu M_{\text{\tiny T}}$  preferably less than 0.25  $\mu M_{\text{\tiny T}}$ 

more preferably less than 0.1  $\mu M$ , and most preferably less than 0.01  $\mu M$ .

Claim 13 (currently amended) The nucleic acid according to any of claims 8 or

9 to 12, wherein the Kd of the nucleic acid is more than 100 nM, preferably more than 10 nM,

more preferably more than 1-nM and most preferably more than 0.05 nM.

Claim 14 (currently amended) The nucleic acid according to any of claim 8 or 9 to 13,

wherein the nucleic acid is of a length selected from the group comprising 15 to 150 nucleotides

 $\underline{\text{in length}}, 20 \text{ to } 100 \text{ nucleotides}, 20 \text{ to } 80 \text{ nucleotides}, 20 \text{ to } 60 \text{ nucleotides}, 20 \text{ to } 50 \text{ nucleotides}$ 

and 30 to 50 nucleotides.

Claim 15 (canceled)

Claim 16 (currently amended) A method for making the generation and/or identification
of a nucleic acid of claim 6 or 8, binding to a target molecule, preferably of a nucleic acid

according to any of claims 6 to 14, comprising the following steps:

a) generating a heterogeneous population of nucleic acids;

b) contacting the population of step a) with the target molecule ghrelin;

c) separating the nucleic acid(s) not interacting with the target molecule ghrelin;

d) optionally separating the nucleic acid(s) interacting with the target molecule ghrelin; and

e) optionally sequencing the nucleic acid(s) interacting with the target molecule.

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characterized in that the target molecule is ghrelin.

Claim 17 (currently amended) The method according to claim 16, <u>further comprising</u> wherein subsequent to step e) a step ea) is carried out, whereby step ea) consists of amplification

of the nucleic acid(s) interacting with the target molecule ghrelin.

Claim 18 (original) The method according to claim 16 or 17, wherein steps b) to d) are

repeated.

Claim 19 (canceled)

Claim 20 (currently amended) A Mmethod for the generating of an making the L-nucleic

acid of claim 10 binding to a target molecule in the natural configuration comprising the

following steps:

generating a heterogeneous population of D-nucleic acids;

b) contacting the population of step a) with an optical antipode of the target

molecule D-ghrelin;

c) separating the D-nucleic acid not interacting with the optical antipode of

the target molecule D-ghrelin;

d) sequencing the D-nucleic acid interacting with the optical antipode of the target

molecule D-ghrelin; and

e) synthesizing the L-nucleic acid sequence(s) which is/are identical to the sequence

of the D-nucleic acid(s) obtained in step d); characterized in that the target molecule is L-

ghrelin and the optical antipode of the target molecule is the D-ghrelin.

Claim 21 (currently amended) The Mmethod according to claim 20 characterized in that

subsequent to step c) the following step is introduced:

ea) <u>further comprising</u> amplifying the D-nucleic acid interacting with the optical

antipode of D-ghrelin the target molecule.

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Claim 22 (currently amended) The Mmethod according to any of claims 20 to claim 20 or 21, characterized in that steps b) to e) are repeated.

Claim 23 (canceled)

Claim 24 (currently amended) Use of a nucleic acid according to any claims 6 to 17 and/or of an antagonist according to any of claims 1 to 5 for the manufacture of a medicament A method of treating a disorder comprising ghrelin or GHSR1a, comprising the step of administering to a patient in need of treatment the nucleic acid of claim 6 or 8, or the antagonist of claim 1 or 2.

Claim 25 (currently amended) The method of Use according to claim 24 characterized in that medicament is for the treatment of a disease or wherein the disorder is selected from the group comprising consisting of obesity[[,]]; the improper regulation of energy balance[[,]]; improper appetite and or body weight[[,]]; cating disorders[[,]]; diabetes[[,]]; improper glucose metabolism[[,]]; tumour[[,]]; improper blood pressure and cardiovascular disease[[s]].

Claim 26 (currently amended)  $\underline{A}$  Composition comprising a <u>the</u> nucleic acid according to any of claim[[s]] 6 or 8 to 14 and/or an <u>the</u> antagonist according to any of claim[[s]] 1 or 2 to 5, and a pharmaceutical acceptable carrier.

Claim 27 (currently amended) A Complex comprising ghrelin and any of the nucleic acid[[s]] according to any of claim[[s]] 6 or 8 to 14, preferably the complex is a crystalline complex.

Claim 28 (canceled)

Claim 29 (currently amended) <u>A</u> Method method for the screening of <u>for</u> a ghrelin antagonist comprising the following steps:

- providing a candidate ghrelin antagonist,
- providing a <u>the</u> nucleic acid according to <u>any of claims claim</u> 6 to 14 or 8, and/or an <u>the</u> antagonist according to <u>any of claims claim</u> 1 to 5 or 2.

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- providing a test system providing a signal in the presence of a ghrelin antagonist,

determining whether the candidate ghrelin antagonist is a ghrelin antagonist.

Claim 30 (currently amended) A Kkit for the detection of ghrelin, comprising [[a]] the nucleic acid according to any of claim[[s]] 6 or 8, to 14 and/or an the antagonist according to any of claim[[s]] 1 or 2 to 5.

Claim 31 (new) The nucleic acid of claim 12, wherein the Kd is less than 0.25 µM.

Claim 32 (new) The nucleic acid of claim 31, wherein the Kd is less than 0.1 µM.

Claim 33 (new) The nucleic acid of claim 32, wherein the Kd is less than 0.01 µM.

Claim 34 (new) The nucleic acid of claim 13, wherein the Kd is more than 10 nM.

Claim 35 (new) The nucleic acid of claim 34, wherein the Kd is more than 1 nM.

Claim 36 (new) The nucleic acid of claim 35, wherein the Kd is more than 0.05 nM.

Claim 37 (new) The nucleic acid of claim 14, wherein the nucleic acid is 20 to 100 nucleotides in length.

Claim 38 (new) The nucleic acid of claim 37, wherein the nucleic acid is 20 to 80 nucleotides in length.

Claim 39 (new) The nucleic acid of claim 38, wherein the nucleic acid is 20 to 60 nucleotides in length.

Claim 40 (new) The nucleic acid of claim 39, wherein the nucleic acid is 20 to 50 nucleotides in length.

Claim 41 (new) The nucleic acid of claim 40, wherein the nucleic acid is 30 to 50 nucleotides in length.

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Claim 42 (new) The nucleic acid of claim 6 comprising the sequence, UAANNCCGAAGGUAACCANUCCUAC, wherein N is any nucleotide.

Claim 43 (new) The nucleic acid of claim 42 comprising the sequence ACNUAANNCCGAAGGUAACCANUCCUACYYACG, wherein Y is no nucleotide or N.

Claim 44 (new) The nucleic acid of claim 43, comprising the sequence AAAACNUAANNCCGAAGGUAACCANUCCUACYYACG.

Claim 45 (new) The complex of claim 27, wherein said complex is crystalline.